Andrew A Dwyer

List of Publications by Year
in descending order

130	8,206	66234	48187
papers	citations	42 h-index	88 g-index

133	133	133	5968
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	European Consensus Statement on congenital hypogonadotropic hypogonadismâ€"pathogenesis, diagnosis and treatment. Nature Reviews Endocrinology, 2015, 11, 547-564.	4.3	664
2	Increasing Insulin Resistance Is Associated with a Decrease in Leydig Cell Testosterone Secretion in Men. Journal of Clinical Endocrinology and Metabolism, 2005, 90, 2636-2641.	1.8	424
3	Relationship Between Testosterone Levels, Insulin Sensitivity, and Mitochondrial Function in Men. Diabetes Care, 2005, 28, 1636-1642.	4.3	392
4	Reversal of Idiopathic Hypogonadotropic Hypogonadism. New England Journal of Medicine, 2007, 357, 863-873.	13.9	362
5	Decreased FGF8 signaling causes deficiency of gonadotropin-releasing hormone in humans and mice. Journal of Clinical Investigation, 2008, 118, 2822-2831.	3.9	348
6	Digenic mutations account for variable phenotypes in idiopathic hypogonadotropic hypogonadism. Journal of Clinical Investigation, 2007, 117, 457-463.	3.9	338
7	Oligogenic basis of isolated gonadotropin-releasing hormone deficiency. Proceedings of the National Academy of Sciences of the United States of America, 2010, 107, 15140-15144.	3.3	313
8	TAC3/TACR3 Mutations Reveal Preferential Activation of Gonadotropin-Releasing Hormone Release by Neurokinin B in Neonatal Life Followed by Reversal in Adulthood. Journal of Clinical Endocrinology and Metabolism, 2010, 95, 2857-2867.	1.8	250
9	Mutations in FGF17, IL17RD, DUSP6, SPRY4, and FLRT3 Are Identified in Individuals with Congenital Hypogonadotropic Hypogonadism. American Journal of Human Genetics, 2013, 92, 725-743.	2.6	227
10	Mutations in fibroblast growth factor receptor 1 cause both Kallmann syndrome and normosmic idiopathic hypogonadotropic hypogonadism. Proceedings of the National Academy of Sciences of the United States of America, 2006, 103, 6281-6286.	3.3	225
11	Predictors of Outcome of Long-Term GnRH Therapy in Men with Idiopathic Hypogonadotropic Hypogonadism. Journal of Clinical Endocrinology and Metabolism, 2002, 87, 4128-4136.	1.8	210
12	Acute Sex Steroid Withdrawal Reduces Insulin Sensitivity in Healthy Men with Idiopathic Hypogonadotropic Hypogonadism. Journal of Clinical Endocrinology and Metabolism, 2007, 92, 4254-4259.	1.8	208
13	Ataxia, Dementia, and Hypogonadotropism Caused by Disordered Ubiquitination. New England Journal of Medicine, 2013, 368, 1992-2003.	13.9	208
14	Mutations inProkineticin 2andProkineticin receptor 2genes in Human Gonadotrophin-Releasing Hormone Deficiency: Molecular Genetics and Clinical Spectrum. Journal of Clinical Endocrinology and Metabolism, 2008, 93, 3551-3559.	1.8	190
15	Mutations in fibroblast growth factor receptor 1 cause Kallmann syndrome with a wide spectrum of reproductive phenotypes. Molecular and Cellular Endocrinology, 2006, 254-255, 60-69.	1.6	176
16	Prioritizing Genetic Testing in Patients With Kallmann Syndrome Using Clinical Phenotypes. Journal of Clinical Endocrinology and Metabolism, 2013, 98, E943-E953.	1.8	157
17	Reversible Kallmann Syndrome, Delayed Puberty, and Isolated Anosmia Occurring in a Single Family with a Mutation in the Fibroblast Growth Factor Receptor 1 Gene. Journal of Clinical Endocrinology and Metabolism, 2005, 90, 1317-1322.	1.8	144
18	Reversal and Relapse of Hypogonadotropic Hypogonadism: Resilience and Fragility of the Reproductive Neuroendocrine System. Journal of Clinical Endocrinology and Metabolism, 2014, 99, 861-870.	1.8	144

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19	Genetic Overlap in Kallmann Syndrome, Combined Pituitary Hormone Deficiency, and Septo-Optic Dysplasia. Journal of Clinical Endocrinology and Metabolism, 2012, 97, E694-E699.	1.8	136
20	GENETICS IN ENDOCRINOLOGY: Genetic counseling for congenital hypogonadotropic hypogonadism and Kallmann syndrome: new challenges in the era of oligogenism and next-generation sequencing. European Journal of Endocrinology, 2018, 178, R55-R80.	1.9	128
21	Genetic basis and variable phenotypic expression of Kallmann syndrome: towards a unifying theory. Trends in Endocrinology and Metabolism, 2011, 22, 249-58.	3.1	127
22	Trial of Recombinant Follicle-Stimulating Hormone Pretreatment for GnRH-Induced Fertility in Patients with Congenital Hypogonadotropic Hypogonadism. Journal of Clinical Endocrinology and Metabolism, 2013, 98, E1790-E1795.	1.8	124
23	Inhibition of Luteinizing Hormone Secretion by Testosterone in Men Requires Aromatization for Its Pituitary But Not Its Hypothalamic Effects: Evidence from the Tandem Study of Normal and Gonadotropin-Releasing Hormone-Deficient Men. Journal of Clinical Endocrinology and Metabolism, 2008, 93, 784-791.	1.8	119
24	Congenital Idiopathic Hypogonadotropic Hypogonadism: Evidence of Defects in the Hypothalamus, Pituitary, and Testes. Journal of Clinical Endocrinology and Metabolism, 2010, 95, 3019-3027.	1.8	115
25	Skeletal Muscle Mitochondria in the Elderly: Effects of Physical Fitness and Exercise Training. Journal of Clinical Endocrinology and Metabolism, 2014, 99, 1852-1861.	1.8	114
26	Expanding the Phenotype and Genotype of Female GnRH Deficiency. Journal of Clinical Endocrinology and Metabolism, 2011, 96, E566-E576.	1.8	97
27	Gonadotrophin replacement for induction of fertility in hypogonadal men. Best Practice and Research in Clinical Endocrinology and Metabolism, 2015, 29, 91-103.	2.2	96
28	The Role of the Prokineticin 2 Pathway in Human Reproduction: Evidence from the Study of Human and Murine Gene Mutations. Endocrine Reviews, 2011, 32, 225-246.	8.9	95
29	Congenital hypogonadotropic hypogonadism and constitutional delay of growth and puberty have distinct genetic architectures. European Journal of Endocrinology, 2018, 178, 377-388.	1.9	95
30	Abrupt decrease in serum testosterone levels after an oral glucose load in men: implications for screening for hypogonadism. Clinical Endocrinology, 2013, 78, 291-296.	1.2	91
31	Human GnRH Deficiency: A Unique Disease Model to Unravel the Ontogeny of GnRH Neurons. Neuroendocrinology, 2010, 92, 81-99.	1.2	87
32	Impaired Fibroblast Growth Factor Receptor 1 Signaling as a Cause of Normosmic Idiopathic Hypogonadotropic Hypogonadism. Journal of Clinical Endocrinology and Metabolism, 2009, 94, 4380-4390.	1.8	82
33	<scp>KLB</scp> , encoding Î²â€Klotho, is mutated in patients with congenital hypogonadotropic hypogonadism. EMBO Molecular Medicine, 2017, 9, 1379-1397.	3.3	77
34	Coding sequence analysis of GNRHR and GPR54 in patients with congenital and adult-onset forms of hypogonadotropic hypogonadism. European Journal of Endocrinology, 2006, 155, S3-S10.	1.9	72
35	MANAGEMENT OF ENDOCRINE DISEASE: Reversible hypogonadotropic hypogonadism. European Journal of Endocrinology, 2016, 174, R267-R274.	1.9	66
36	Identifying the unmet health needs of patients with congenital hypogonadotropic hypogonadism using a web-based needs assessment: implications for online interventions and peer-to-peer support. Orphanet Journal of Rare Diseases, 2014, 9, 83.	1.2	63

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37	GnRH-Deficient Phenotypes in Humans and Mice with Heterozygous Variants inKISS1/Kiss1. Journal of Clinical Endocrinology and Metabolism, 2011, 96, E1771-E1781.	1.8	59
38	TRANSITION IN ENDOCRINOLOGY: Hypogonadism in adolescence. European Journal of Endocrinology, 2015, 173, R15-R24.	1.9	59
39	The Relative Role of Gonadal Sex Steroids and Gonadotropin-Releasing Hormone Pulse Frequency in the Regulation of Follicle-Stimulating Hormone Secretion in Men. Journal of Clinical Endocrinology and Metabolism, 2008, 93, 2686-2692.	1.8	55
40	Congenital hypogonadotropic hypogonadism with split hand/foot malformation: a clinical entity with a high frequency of FGFR1 mutations. Genetics in Medicine, 2015, 17, 651-659.	1.1	55
41	International practice settings, interventions and outcomes of nurse practitioners in geriatric care: A scoping review. International Journal of Nursing Studies, 2018, 78, 61-75.	2.5	54
42	The Long-Term Clinical Follow-Up and Natural History of Men with Adult-Onset Idiopathic Hypogonadotropic Hypogonadism. Journal of Clinical Endocrinology and Metabolism, 2010, 95, 4235-4243.	1.8	45
43	DCC/NTN1 complex mutations in patients with congenital hypogonadotropic hypogonadism impair GnRH neuron development. Human Molecular Genetics, 2018, 27, 359-372.	1.4	42
44	Î²-Klotho deficiency protects against obesity through a crosstalk between liver, microbiota, and brown adipose tissue. JCI Insight, 2017, 2, .	2.3	41
45	Relative Roles of Inhibin B and Sex Steroids in the Negative Feedback Regulation of Follicle-Stimulating Hormone in Men across the Full Spectrum of Seminiferous Epithelium Function. Journal of Clinical Endocrinology and Metabolism, 2008, 93, 1809-1814.	1.8	40
46	Neuron-Derived Neurotrophic Factor Is Mutated in Congenital Hypogonadotropic Hypogonadism. American Journal of Human Genetics, 2020, 106, 58-70.	2.6	39
47	Evaluating CHARGE syndrome in congenital hypogonadotropic hypogonadism patients harboring CHD7 variants. Genetics in Medicine, 2018, 20, 872-881.	1.1	38
48	Metaâ€analysis of the effectiveness of nursing discharge planning interventions for older inpatients discharged home. Journal of Advanced Nursing, 2018, 74, 788-799.	1.5	37
49	Testosterone restoration using enclomiphene citrate in men with secondary hypogonadism: a pharmacodynamic and pharmacokinetic study. BJU International, 2013, 112, 1188-1200.	1.3	35
50	Psychosexual Development in Men with Congenital Hypogonadotropic Hypogonadism on Long-Term Treatment: A Mixed Methods Study. Sexual Medicine, 2015, 3, 32-41.	0.9	34
51	Adherence to treatment in men with hypogonadotrophic hypogonadism. Clinical Endocrinology, 2017, 86, 377-383.	1.2	32
52	Fertility induction in hypogonadotropic hypogonadal men. Clinical Endocrinology, 2018, 89, 712-718.	1.2	32
53	An ancient founder mutation in PROKR2 impairs human reproduction. Human Molecular Genetics, 2012, 21, 4314-4324.	1.4	31
54	Beyond hormone replacement: quality of life in women with congenital hypogonadotropic hypogonadism. Endocrine Connections, 2017, 6, 404-412.	0.8	31

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55	Functional Hypogonadotropic Hypogonadism in Men: Underlying Neuroendocrine Mechanisms and Natural History. Journal of Clinical Endocrinology and Metabolism, 2019, 104, 3403-3414.	1.8	28
56	Multidisciplinary management of diabetic kidney disease. JBI Database of Systematic Reviews and Implementation Reports, 2016, 14, 169-207.	1.7	26
57	Developing and evaluating rare disease educational materials co-created by expert clinicians and patients: the paradigm of congenital hypogonadotropic hypogonadism. Orphanet Journal of Rare Diseases, 2017, 12, 57.	1.2	26
58	Psychological Aspects of Congenital Hypogonadotropic Hypogonadism. Frontiers in Endocrinology, 2019, 10, 353.	1.5	26
59	Symptom perception in heart failure: a scoping review on definition, factors and instruments. European Journal of Cardiovascular Nursing, 2020, 19, 100-117.	0.4	25
60	Role of fibroblast growth factor (FGF) signaling in the neuroendocrine control of human reproduction. Molecular and Cellular Endocrinology, 2011, 346, 37-43.	1.6	24
61	Hormonal control of spermatogenesis in men: Therapeutic aspects in hypogonadotropic hypogonadism. Annales D'Endocrinologie, 2014, 75, 98-100.	0.6	21
62	Effectiveness of nursing discharge planning interventions on health-related outcomes in discharged elderly inpatients. JBI Database of Systematic Reviews and Implementation Reports, 2016, 14, 217-260.	1.7	21
63	Society for Endocrinology <scp>UK</scp> guidance on the evaluation of suspected disorders of sexual development: emphasizing the opportunity to predict adolescent pubertal failure through a neonatal diagnosis of absent minipuberty. Clinical Endocrinology, 2017, 86, 305-306.	1.2	21
64	Congenital hypogonadotropic hypogonadism/Kallmann syndrome is associated with statural gain in both men and women: a monocentric study. European Journal of Endocrinology, 2020, 182, 185.	1.9	21
65	Accuracy, satisfaction and usability of a flash glucose monitoring system among children and adolescents with type 1 diabetes attending a summer camp. Pediatric Diabetes, 2018, 19, 1276-1284.	1.2	20
66	Anatomy and Physiology of theÂHypothalamic-Pituitary-Gonadal (HPG) Axis. , 2019, , 839-852.		20
67	Congenital Hypogonadotropic Hypogonadism with Anosmia and Gorlin Features Caused by a PTCH1 Mutation Reveals a New Candidate Gene for Kallmann Syndrome. Neuroendocrinology, 2021, 111, 99-114.	1.2	20
68	Testis morphology in patients with idiopathic hypogonadotropic hypogonadism. Human Reproduction, 2006, 21, 1033-1040.	0.4	19
69	Comparative functional analysis of two fibroblast growth factor receptor 1 (FGFR1) mutations affecting the same residue (R254W and R254Q) in isolated hypogonadotropic hypogonadism (IHH). Gene, 2013, 516, 146-151.	1.0	19
70	The Metabolic Syndrome in Central Hypogonadotrophic Hypogonadism. Frontiers of Hormone Research, 2018, 49, 156-169.	1.0	19
71	Evaluating coâ€created patientâ€facing materials to increase understanding of genetic test results. Journal of Genetic Counseling, 2021, 30, 598-605.	0.9	19
72	Acceptance of the Advanced Practice Nurse in Lung Cancer Role by Healthcare Professionals and Patients: A Qualitative Exploration. Journal of Nursing Scholarship, 2018, 50, 540-548.	1.1	18

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73	Patient Perspectives on Nurse-led Consultations Within a Pilot Structured Transition Program for Young Adults Moving From an Academic Tertiary Setting to Community-based Type 1 Diabetes Care. Journal of Pediatric Nursing, 2018, 38, 99-105.	0.7	17
74	Symptom perception in heart failure â€“ Interventions and outcomes: A scoping review. International Journal of Nursing Studies, 2021, 116, 103524.	2.5	17
75	Impact of Acute Biochemical Castration on Insulin Sensitivity in Healthy Adult Men. Endocrine Research, 2010, 35, 71-84.	0.6	15
76	Genetic testing facilitates prepubertal diagnosis of congenital hypogonadotropic hypogonadism. Clinical Genetics, 2017, 92, 213-216.	1.0	14
77	Feasibility of advanced practice nursing in lung cancer consultations during early treatment: A phase II study. European Journal of Oncology Nursing, 2017, 29, 106-114.	0.9	14
78	Psychosexual effects resulting from delayed, incomplete, or absent puberty. Current Opinion in Endocrine and Metabolic Research, 2020, 14, 15-21.	0.6	14
79	GnRH stimulation testing and serum inhibin B in males: insufficient specificity for discriminating between congenital hypogonadotropic hypogonadism from constitutional delay of growth and puberty. Human Reproduction, 2020, 35, 2312-2322.	0.4	13
80	Natural History of Growth Hormone Deficiency in a Pediatric Cohort. Hormone Research in Paediatrics, 2015, 83, 252-261.	0.8	12
81	Thigh and abdominal adipose tissue depot associations with testosterone levels in postmenopausal females. Clinical Endocrinology, 2019, 90, 433-439.	1.2	12
82	Congenital hypogonadotropic hypogonadism: implications of absent mini-puberty. Minerva Endocrinologica, 2016, 41, 188-95.	1.7	12
83	Parent of Origin Effects on Family Communication of Risk in BRCA+ Women: A Qualitative Investigation of Human Factors in Cascade Screening. Cancers, 2020, 12, 2316.	1.7	11
84	A sexual health course for advanced practice registered nurses: Effect on preparedness, comfort, and confidence in delivering comprehensive care. Nurse Education Today, 2020, 92, 104506.	1.4	10
85	Role of Seminiferous Tubular Development in Determining the FSH versus LH Responsiveness to GnRH in Early Sexual Maturation. Neuroendocrinology, 2009, 90, 260-268.	1.2	9
86	Patient and healthcare professional eHealth literacy and needs for systemic sclerosis support: a mixed methods study. RMD Open, 2021, 7, e001783.	1.8	9
87	Absence of Central Circadian Pacemaker Abnormalities in Humans With Loss of Function Mutation in Prokineticin 2. Journal of Clinical Endocrinology and Metabolism, 2014, 99, E561-E566.	1.8	8
88	Parent of origin differences in psychosocial burden and approach to BRCA risk management. Breast Journal, 2020, 26, 734-738.	0.4	7
89	Insight Into the Ontogeny of GnRH Neurons From Patients Born Without a Nose. Journal of Clinical Endocrinology and Metabolism, 2020, 105, 1538-1551.	1.8	7
90	The lack of effect of insulin on luteinizing hormone pulsatility in healthy male volunteers provides evidence of a sexual dimorphism in the metabolic regulation of reproductive hormones. American Journal of Clinical Nutrition, 2012, 96, 283-288.	2.2	6

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91	Transition of Care from Childhood to Adulthood: Congenital Hypogonadotropic Hypogonadism. Endocrine Development, 2018, 33, 82-98.	1.3	6
92	Non-invasive assessment of coronary endothelial function in children and adolescents with type 1 diabetes mellitus using isometric handgrip exerciseâ€"MRI: A feasibility study. PLoS ONE, 2020, 15, e0228569.	1.1	5
93	Insulin-like Growth Factor 1, but Not Insulin-Like Growth Factor-Binding Protein 3, Predicts Central Precocious Puberty in Girls 6â€"8 Years Old: A Retrospective Study. Hormone Research in Paediatrics, 2021, 94, 1-8.	0.8	5
94	Coping response and family communication of cancer risk in men harboring aBRCAmutation: A mixed methods study. Psycho-Oncology, 2022, 31, 486-495.	1.0	5
95	A novel CHD7 mutation in an adolescent presenting with growth and pubertal delay. Annals of Pediatric Endocrinology and Metabolism, 2019, 24, 49-54.	0.8	5
96	Acute Stress Masking the Biochemical Phenotype of Partial Androgen Insensitivity Syndrome in a Patient with a Novel Mutation in the Androgen Receptor. Journal of Clinical Endocrinology and Metabolism, 2004, 89, 1053-1058.	1.8	4
97	European-South Africa collaboration on the genetic basis of gonadotropin-releasing hormone deficiency in failure to progress through puberty and infertility. Journal of Endocrinology Metabolism and Diabetes of South Africa, 2015, 20, 67-68.	0.4	4
98	Developing a rare disease chronic care model: Management of systemic sclerosis (MANOSS) study protocol. Journal of Advanced Nursing, 2019, 75, 3774-3791.	1.5	4
99	Validating online approaches for rare disease research using latent class mixture modeling. Orphanet Journal of Rare Diseases, 2021, 16, 209.	1.2	4
100	Effectiveness of nursing discharge planning interventions on health-related outcomes in elderly inpatients discharged home: a systematic review protocol. JBI Database of Systematic Reviews and Implementation Reports, 2013, 11, 1-12.	1.7	4
101	Navigating Disrupted Puberty: Development and Evaluation of a Mobile-Health Transition Passport for Klinefelter Syndrome. Frontiers in Endocrinology, 0, 13, .	1.5	4
102	Patient perceptions of peripheral artery disease: A cross-sectional study of hospitalized adults. Journal of Vascular Nursing, 2019, 37, 188-193.	0.2	3
103	Framing Effects on Decision-Making for Diagnostic Genetic Testing: Results from a Randomized Trial. Genes, 2021, 12, 941.	1.0	3
104	Endocrine Nurses Society Position Statement on Transgender and Gender Diverse Care. Journal of the Endocrine Society, 2021, 5, bvab105.	0.1	3
105	Comment on reversal of hypogonadotropic hypogonadism in a Chinese cohort. Asian Journal of Andrology, 2015, 17, 508.	0.8	3
106	Exploring Rare Disease Patient Attitudes and Beliefs regarding Genetic Testing: Implications for Person-Centered Care. Journal of Personalized Medicine, 2022, 12, 477.	1.1	3
107	Genetic Competencies for Effective Pediatric Endocrine Nursing Practice. Journal of Pediatric Nursing, 2019, 48, 127-128.	0.7	2
108	Classification of Hypothalamic-Pituitary-Gonadal (HPG) Axis Endocrine Disorders. , 2019, , 853-870.		2

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109	Congenital hyperinsulinism: 2 case reports with different rare variants in ABCC8. Annals of Pediatric Endocrinology and Metabolism, 2021, 26, 60-65.	0.8	2
110	A rare disease patient-reported outcome measure: revision and validation of the German version of the Systemic Sclerosis Quality of Life Questionnaire (SScQoL) using the Rasch model. Orphanet Journal of Rare Diseases, 2021, 16, 356.	1.2	2
111	Impact of BRCA Status on Reproductive Decision-Making and Self-Concept: A Mixed-Methods Study Informing the Development of Tailored Interventions. Cancers, 2022, 14, 1494.	1.7	2
112	Correspondence on â€œEnsuring best practice in genomics education and evaluation: Reporting item standards for education and its evaluation in genomics (RISE2 Genomics)â€•by Nisselle etÂal. Genetics in Medicine, 2022, 24, 962-963.	1.1	2
113	PENS Position Statement on Bullying Prevention. Journal of Pediatric Nursing, 2018, 39, 91-93.	0.7	1
114	Minipuberty: A Primer for Pediatric Nurses. Journal of Pediatric Nursing, 2020, 50, 138-139.	0.7	1
115	What's missing in sex chromosome aneuploidies? Representation and inclusion. Journal of Pediatric Nursing, 2022, 62, 202-204.	0.7	1
116	A Comparison of the Blood Glucose, Growth Hormone, and Cortisol Responses to Two Doses of Insulin (0.15â€‰U/kg vs. 0.10â€‰U/kg) in the Insulin Tolerance Test: A Single-Centre Audit of 174 Cases. International Journal of Endocrinology, 2022, 2022, 1-8.	0.6	1
117	TAC3/TACR3 Mutations Reveal Preferential Activation of GnRH Release by Neurokinin B in Neonatal Life Followed by Reversal in Adulthood. Endocrinology, 2010, 151, 1970-1971.	1.4	0
118	GnRH-Deficient Phenotypes in Humans and Mice With Heterozygous Variants in KISS1/Kiss1. Obstetrical and Gynecological Survey, 2012, 67, 546-547.	0.2	0
119	004-Caring Matters: A Cross-Sectional Study of Patient Perspectives on Nurse-led Consultations as Part of Structured Transitional Care for Type 1 Diabetes. Journal of Pediatric Nursing, 2017, 37, 144.	0.7	0
120	Fertility and the Hypogonadal Male. , 2019, , 94-105.		0
121	Spermatogenesis and Assisted Fertility Treatment. , 2019, , 903-923.		0
122	Evaluation of Endocrine Disorders of the Hypothalamic-Pituitary-Gonadal (HPG) Axis. , 2019, , 871-883.		0
123	Editorial: New Aspects in Hypogonadism. Frontiers in Endocrinology, 2020, 11, 426.	1.5	0
124	Adherence to treatment for chronic hypogonadism: the role of illness perceptions and depressive symptoms. Endocrine Abstracts, 0, , .	0.0	0
125	Research topic: identifying the needs of patients with Congenital Hypogonadotrophic Hypogonadism, implications for nursing practice. Endocrine Abstracts, 0, , .	0.0	0
126	Unmet health and information needs of women with hypogonadotropic hypogonadism. Endocrine Abstracts, 0, , .	0.0	0

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127	Secondary Infertility. Endocrine Abstracts, 0, , .	0.0	0
128	Clinical practice overlap and seamless care - links between hypogonadism, the metabolic syndrome and type 2 diabetes. Endocrine Abstracts, 0, , .	0.0	0
129	Transitional Care in Endocrinology. , 2020, , 281-317.		0
130	A Developmental Perspective Sheds Light on Reproductive Differences Between Congenital and Acquired Hypogonadism. Journal of Clinical Endocrinology and Metabolism, 2022, 107, e3955-e3956.	1.8	0